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## IN THE HOUSE OF REPRESENTATIVES

## HOUSE BILL NO. 208

## BY HEALTH AND WELFARE COMMITTEE

	BI HEALIH AND WELFARE COMMITTEE
1 2 3 4	AN ACT RELATING TO PHARMACISTS; AMENDING SECTION 54-1704, IDAHO CODE, TO REMOVE A PROVISION REGARDING FEDERAL FOOD AND DRUG ADMINISTRATION-APPROVED LA- BELING.
5	Be It Enacted by the Legislature of the State of Idaho:
6 7	SECTION 1. That Section 54-1704, Idaho Code, be, and the same is hereby amended to read as follows:
8 9 10	54-1704. PRACTICE OF PHARMACY. "Practice of pharmacy" means: (1) The interpretation, evaluation and dispensing of prescription drug orders;
11 12 13	(2) Participation in drug and device selection, drug administration, prospective and retrospective drug reviews and drug or drug-related research;
14 15 16	<ul><li>(3) The provision of patient counseling and the provision of those acts or services necessary to provide pharmaceutical care;</li><li>(4) The responsibility for:</li></ul>
17 18 19	(a) Compounding and labeling of drugs and devices, except labeling by a manufacturer, repackager or distributor of nonprescription drugs and commercially packaged legend drugs and devices;
19 20 21	(b) Proper and safe storage of drugs and devices, and maintenance of proper records for them; and
22 23 24	(c) The offering or performing of those acts, services, operations or transactions necessary to the conduct, operation, management and control of pharmacy;
25 26 27	<ul><li>(5) The prescribing of:</li><li>(a) Agents for active immunization when prescribed for susceptible persons six (6) years of age or older for the protection from communica-</li></ul>
28 29	ble disease; and (b) Drugs, drug categories, or devices that are prescribed in accor-
30 31	dance with the product's federal food and drug administration-approved labeling and that are limited to conditions that:
32 33	<ul><li>(i) Do not require a new diagnosis;</li><li>(ii) Are minor and generally self-limiting;</li></ul>
34 35 36	(iii) Have a test that is used to guide diagnosis or clinical decision-making and are waived under the federal clinical laboratory improvement amendments of 1988; or

(iv) In the professional judgment of the pharmacist, threaten

the health or safety of the patient should the prescription not be

immediately dispensed. In such cases, only sufficient quantity

may be provided until the patient is able to be seen by another

The board shall not adopt any rules authorizing a pharmacist to prescribe a controlled drug, compounded drug or biological product.